

**Amendments to the Claims:**

Please cancel claims 3, 10-18, 20, and 28-30 without disclaimer or prejudice to applicants' right to pursue the subject matters of these claims in the future.

Pursuant to 37 C.F.R. §1.121(c), this listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims:**

1. (Currently Amended) A method of treating a fibrosis-related pathology in a subject which comprises administering to the subject a therapeutically effective amount of a pharmaceutical composition comprising a Phospholipase D<sub>3</sub> inhibitor so as to thereby treat the subject.
2. (Currently Amended) The method of claim 1 wherein the pharmaceutical composition comprises an oligoribonucleotide or oligonucleotide which down-regulates the expression of gene Phospholipase D<sub>3</sub> by at least 50% as compared to a control.
3. (Canceled)
4. (Currently Amended) The method of claim 1 wherein the Phospholipase D<sub>3</sub> inhibitor is an antisense oligonucleotide.
5. (Currently Amended) The method of claim 1 wherein the Phospholipase D<sub>3</sub> inhibitor is a Phospholipase D<sub>3</sub> siRNA.
6. (Currently Amended) The method of claim 1 wherein the Phospholipase D<sub>3</sub> inhibitor is an expression vector comprising a nucleic acid molecule encoding Phospholipase D<sub>3</sub> siRNA.

7. (Currently Amended) The method of claim 1 wherein the Phospholipase D<sub>3</sub> inhibitor is an antibody which binds specifically to Phospholipase D<sub>3</sub> polypeptide.
8. (Currently Amended) The method of claim 1 wherein the fibrosis-related pathology is chronic renal insufficiency, chronic renal insufficiency, nephropathy, liver fibrosis, cardiac fibrosis or kidney fibrosis.
9. (Original) The method of claim 1 wherein the fibrosis-related pathology is ocular scarring or cataract.
- 10-18. (Canceled)
19. (Currently Amended) A pharmaceutical composition for the treatment of fibrosis comprising as an active ingredient a Phospholipase D<sub>3</sub> inhibitor together with a pharmaceutically acceptable carrier.
20. (Canceled)
21. (Currently Amended) The pharmaceutical composition of claim 19 wherein the Phospholipase D<sub>3</sub> inhibitor is an oligoribonucleotide or oligonucleotide which down-regulates the expression of gene Phospholipase D<sub>3</sub> by at least 50% as compared to a control.
22. (Currently Amended) The pharmaceutical composition of claim 19 wherein the Phospholipase D<sub>3</sub> inhibitor is an antisense oligonucleotide.
23. (Currently Amended) The pharmaceutical composition of claim 19 wherein the Phospholipase D<sub>3</sub> inhibitor is a Phospholipase D<sub>3</sub> siRNA.
24. (Currently Amended) The pharmaceutical composition of claim 19 wherein the Phospholipase D<sub>3</sub> inhibitor is an expression vector comprising a nucleic acid molecule encoding Phospholipase D<sub>3</sub> siRNA.

Applicants: Orna Mor, et al.  
U.S. Serial No.: Not Yet Known  
Filed: Herewith  
Page 7

25. (Currently Amended) The pharmaceutical composition of claim 19 wherein the Phospholipase D<sub>3</sub> inhibitor is an antibody which binds specifically to Phospholipase D<sub>3</sub> polypeptide.
26. (Currently Amended) The pharmaceutical composition of claim 19 wherein the fibrosis-related pathology is chronic renal insufficiency, chronic renal insufficiency, nephropathy, liver fibrosis, cardiac fibrosis or kidney fibrosis.
27. (Original) The pharmaceutical composition of claim 19 wherein the fibrosis-related pathology is ocular scarring or cataract.
- 28-30. (Canceled)